

about polysaccharide selected from xanthan gum and hydroxypropylmethylcellulose (HPMC) as a therapeutically active agent in an amount effective to treat inflammatory bowel disease, together with a pharmaceutically acceptable carrier or vehicle.

2. (Amended) [A] The composition [as claimed in] according to Claim 1, wherein the polysaccharide is xanthan gum.

3. (Amended) [A] The composition [as claimed in] according to Claim 1, wherein the polysaccharide is HPMC.

4. (Amended) [A] The composition [as claimed in] according to [any one of the preceding claims] Claim 1, wherein the polysaccharide is present as the sole therapeutically active ingredient.

5. (Amended) [A] The [DRO] composition [as claimed in] according to [any one of the preceding claims] Claim 1 (which is a DRO composition).

6. (Amended) [A] The [DRO] composition [as claimed in] according to Claim 5 which DRO composition is an enteric coated dosage form adapted to release its contents within the region of the jejunum to the colon.

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7. (Amended) [A] ~~The rectally administrable composition [as claimed in]~~
according to [any one of Claims 1 to 4] Claim 1.

8. (Amended) [A] ~~The rectally administrable composition [as claimed in]~~
according to Claim 7 which is a liquid enema or foam enema.

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9. (Amended) [A] ~~The [liquid enema] composition [as claimed in]~~ according to
Claim [8] 2, which is a liquid enema containing [wherein the polysaccharide is] xanthan
gum in a concentration of about 0.4 to about 2% w/w (based on the composition).

10. (Amended) [A] ~~The [foam enema] composition [as claimed in]~~ according to
Claim [8] 2, which is a foam enema containing [wherein the polysaccharide is] xanthan
gum in a concentration of about 1.4 to about 2.5 % w/w (based on the composition).

11. (Amended) [A] ~~The [liquid enema] composition [as claimed in]~~ according to
Claim [8] 3, which is a liquid enema containing [wherein the polysaccharide is] HPMC in
a concentration of about 1 to about 20 % w/w (based on the composition).

12. (Amended) [A] ~~The [foam enema] composition [as claimed in]~~ according to
Claim [8] 3, which is a foam enema containing [wherein the polysaccharide is] HPMC in
a concentration of about 2.5 to about 25% w/w (based on the composition).

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13. (Amended) [A] The rectally administrable composition [as claimed in] according to Claim 7 [or Claim 8], wherein the polysaccharide is xanthan gum in an amount of about 400 to about 2000 mg per unit dose.

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14. (Amended) [A] The rectally administrable composition [as claimed in] according to Claim 7 [or Claim 8], wherein the polysaccharide is HPMC in an amount of about 1 to about 20 g per unit dose.

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15. (Amended) [A] The DRO composition [as claimed in] according to Claim 5 [or Claim 6, wherein the] in unit dose form containing about 400 to about 2000 mg of the polysaccharide [is 400 to 2000 mg] per unit dose.

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22. (Amended) A method for the treatment or prophylaxis of inflammatory bowel disease (IBD) comprising contacting the diseased mucosa of the gastro-intestinal tract with a therapeutic amount of a polysaccharide selected from xanthan gum and hydroxypropylmethylcellulose (HPMC).

Add the following new claims:

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- - 23. The liquid enema according to Claim 11, wherein the HPMC is in a concentration of 5 to 20 % w/w (based on the composition).

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24. The method according to Claim 22 wherein the disease state is pouchitis.

25. The method according to Claim 22 wherein the disease state is left-sided ulcerative colitis.

26. The method according to Claim 22 wherein the disease state is Crohn's disease.

27. A liquid enema for the treatment or prophylaxis of inflammatory bowel disease (IBD) comprising xanthan gum in a concentration of about 0.4 to about 2 % w/w (based on the composition) as a therapeutically active agent in an amount effective to treat inflammatory bowel disease, together with a pharmaceutically acceptable carrier or vehicle.

28. A foam enema for the treatment or prophylaxis of inflammatory bowel disease (IBD) comprising xanthan gum in a concentration of about 1.4 to 2.5 % w/w (based on the composition) as a therapeutically active agent in an amount effective to treat inflammatory bowel disease, together with a pharmaceutically acceptable carrier or vehicle. - -